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Complete if Known

Application Number 10/625,124

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First Named Inventor Dobrowolski, Steven

Art Unit 1645

STATEMENT BY APPLICANT

| Comparison of the comp

NON PATENT LITERATURE DOCUMENTS						
Examiner Initials*	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.				
Gn		POMPONIO, RJ et al., Profound biotinidase deficiency caused by a point mutation that creates, Human Molecular Genetics (1997), 6(5), 739-745.				
Cay	COLE, H et al., Human Serum Biotinidase: cDNA Cloning, Sequence, and Characterization, J. Biol. Chem. (1994), 269(9), 6566-6570.					
Or		HEARD, Gregory et al., A Screening Method for Biotinidase Deficiency in Newborns, Clin. Chem. (1984), 30/1, 125-127.				

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